Amendments to the Claims:

This listing of claims will replace all prior versions, and listings of claims in the application:

Listing of Claims:

- 1. 63. (Canceled)
- 64. (Withdrawn- currently amended) The fragment complementation system of claim 63 78, wherein said first oligopeptide and said second oligopeptide comprise a signal peptide that translocates said first oligopeptide and said second oligopeptide through the plasma membrane of a host cell in which said first oligopeptide and said second oligopeptide are expressed.
 - 65. (Canceled)
- 66. (Currently amended) The fragment complementation system of claim 63
 79, wherein said Class A <u>TEM-1</u> β-lactamase protein comprises amino acids 26 to 288 of the following sequence:

His Pro Glu Thr Leu Val Lys Val Lys Asp Ala Glu Asp Gln Leu Gly 30 35 40 26 Ala Arg Val Gly Tyr Ile Glu Leu Asp Leu Asn Ser Gly Lys Ile Leu 45 50 55 Glu Ser Phe Arg Pro Glu Glu Arg Phe Pro Met Met Ser Thr Phe Lys 65 60 70 Val Leu Leu Cys Gly Ala Val Leu Ser Arg Ile Asp Ala Gly Gln Glu 75 85 80 Gln Leu Gly Arg Arg Ile His Tyr Ser Gln Asn Asp Leu Val Glu Tyr 90 95 100 105 Ser Pro Val Thr Glu Lys His Leu Thr Asp Gly Met Thr Val Arg Glu

115

120

110

Leu Cys Ser Ala Ala Ile Thr Met Ser Asp Asn Thr Ala Ala Asn Leu Leu Leu Thr Thr Ile Gly Gly Pro Lys Glu Leu Thr Ala Phe Leu His Asn Met Gly Asp His Val Thr Arg Leu Asp Arg Trp Glu Pro Glu Leu Asn Glu Ala Ile Pro Asn Asp Glu Arg Asp Thr Thr Met Pro Val Ala Met Ala Thr Thr Leu Arg Lys Leu Leu Thr Gly Glu Leu Leu Thr Leu Ala Ser Arg Gln Gln Leu Ile Asp Trp Met Glu Ala Asp Lys Val Ala Gly Pro Leu Leu Arg Ser Ala Leu Pro Ala Gly Trp Phe Ile Ala Asp Lys Ser Gly Ala Gly Glu Arg Gly Ser Arg Gly Ile Ile Ala Ala Leu Gly Pro Asp Gly Lys Pro Ser Arg Ile Val Val Ile Tyr Thr Thr Gly Ser Gln Ala Thr Met Asp Glu Arg Asn Arg Gln Ile Ala Glu Ile Gly Ala Ser Leu Ile Lys His Trp (SEQ ID NO:2);

67. (Canceled).

and Leu 198.

wherein said first and second break-point is between amino acid residues Glu 197

Appl. No. 10/668,778 Amdt. dated November 11, 2008 Reply to Office Action of June 12, 2008

- 68. (Withdrawn-Currently amended) The fragment complementation system of claim 63 78, wherein said fragment complementation system further comprises a first peptide that enhances the functional reconstitution of said N-terminal fragment and said C-terminal fragment in comparison with the identical system without said first peptide, wherein said first peptide is 3-12 amino acids in length.
- 69. (Withdrawn-Previously Presented) The fragment complementation system of claim 68, wherein said first peptide is 3 amino acids in length.
- 70. (Withdrawn-Previously Presented) The fragment complementation system of claim 69, wherein said first peptide is covalently bonded to the active site of a thioredoxin protein, wherein the sequence of said first peptide is GRE.
- 71. (Currently amended) The fragment complementation system of claim 63

 78, wherein

 said first polypeptide linker is 3-30 amino acids in length; and wherein said second polypeptide linker is 3-30 amino acids in length.
- 72. (Currently amended) The fragment complementation system of claim 71, further comprising a first complementation enhancement peptide fused between the N-terminal fragment of the Class A <u>TEM-1</u> β-lactamase protein and the first polypeptide linker; and

a second complementation enhancement peptide fused between the C-terminal fragment of the Class A TEM-1 β -lactamase protein and the second polypeptide linker.

73. (Previously Presented) The fragment complementation system of claim 72, wherein

the sequence of said first complementation enhancement peptide is selected from the group consisting of HSE, GRE, EKR, and NGR, and

the sequence of said second complementation enhancement peptide is selected from the group consisting of REQ, QGN, DGR, GRR and GNS.

Appl. No. 10/668,778 Amdt. dated November 11, 2008 Reply to Office Action of June 12, 2008

74. (Previously presented) The fragment complementation system of claim 73, wherein

the sequence of said first complementation enhancement peptide is HSE, and the sequence of said second complementation enhancement peptide is REQ.

- 75. (Previously presented) The fragment complementation system of claim 73, wherein the sequence of said first complementation enhancement peptide is NGR, and the sequence of said second complementation enhancement peptide is QGN or GNS.
- 76. (Previously presented) The fragment complementation system of claim 73, wherein the sequence of said first complementation enhancement peptide is GRE, and the sequence of said second complementation enhancement peptide is DGR.
- 77. (Previously presented) The fragment complementation system of claim 73, wherein the sequence of said first complementation enhancement peptide is EKR, and the sequence of said second complementation enhancement peptide is GRR.
- 78 (New) A fragment complementation system, said system comprising a first oligopeptide sequence and a second oligopeptide sequence;

wherein said first oligopeptide sequence is a fusion protein comprising, in the direction of translation, an N-terminal fragment of a TEM-1 β -lactamase protein, not less than 25 amino acids in length, fused through a C-terminal residue to a first flexible polypeptide linker and a first interactor domain; and

wherein said second oligopeptide sequence is a fusion protein comprising, in the direction of translation, a second interactor domain, a second flexible polypeptide linker fused through an N-terminal residue to a C-terminal fragment of the TEM-1 β-lactamase protein, not less than 25 amino acids in length;

wherein said C-terminal residue and N-terminal residue are located in a solvent exposed loop between amino acid residues Thr 195 and Ala 202 of the TEM-1 β -lactamase protein and,

Appl. No. 10/668,778 Amdt. dated November 11, 2008 Reply to Office Action of June 12, 2008

wherein upon binding of said first interactor domain with said second interactor domain, said N-terminal fragment and said C-terminal fragment reconstitute to form a functional TEM-1 β -lactamase protein.

79. (New) The fragment complementation system of claim 78, wherein the C-terminal residue of the N-terminal fragment is Glu 197, and the C-terminal residue of the N-terminal fragment is Leu 198.